UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/590,492	06/01/2007	Takahide Kohro	032217A	1086	
	7590 05/15/200 , HATTORI, DANIEL		EXAMINER		
1250 CONNEC	1250 CONNECTICUT AVENUE, NW			RAE, CHARLESWORTH E	
SUITE 700 WASHINGTON, DC 20036			ART UNIT	PAPER NUMBER	
			1611		
			MAIL DATE	DELIVERY MODE	
			05/15/2008	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)
	10/590,492	KOHRO ET AL.
Office Action Summary	Examiner	Art Unit
	CHARLESWORTH RAE	1611
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with the c	correspondence address
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D - Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period - Failure to reply within the set or extended period for reply will, by statute Any reply received by the Office later than three months after the mailin earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 136(a). In no event, however, may a reply be tin will apply and will expire SIX (6) MONTHS from e, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. D (35 U.S.C. § 133).
Status		
1) Responsive to communication(s) filed on <u>24 A</u> 2a) This action is FINAL . 2b) This 3) Since this application is in condition for alloware closed in accordance with the practice under B	s action is non-final. nce except for formal matters, pro	
Disposition of Claims		
 4) Claim(s) 1-15 is/are pending in the application 4a) Of the above claim(s) is/are withdra 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) 1-15 are subject to restriction and/or 	wn from consideration.	
Application Papers		
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) accomposed applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Examine 11.	cepted or b) objected to by the I drawing(s) be held in abeyance. See tion is required if the drawing(s) is objected to by the I	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) ☐ Acknowledgment is made of a claim for foreign a) ☐ All b) ☐ Some * c) ☐ None of: 1. ☐ Certified copies of the priority document 2. ☐ Certified copies of the priority document 3. ☐ Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	ts have been received. ts have been received in Applicati rity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal F 6) Other:	ate

DETAILED ACTION

Claims 1-15 are currently pending in this application and are the subject of the Office action.

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

It is noted that the "use claims" (i.e. 4, 5, 6, and 11) have been considered as a separate group because it is unclear whether these claims are directed to a method of using or method of making.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a <u>single</u> invention to which the claims must be restricted.

- I. Claims 1-3, drawn to a nuclear transfer promoter for Cdc42 protein comprising an isopremoid synthesis inhibitor and/or a geranylgeranyl transferase inhibitor. If this Group is elected, then the below Summarized Species Election is also required.
- II. Claims 7, drawn to a method of promoting the transfer of Cd42 protein into a nucleus, which comprises administering an isoprenoid synthesis inhibitor and/or a geranylgeranyl transferase inhibitor to a cell. If this Group is elected, then the below Summarized Species Election is also required.

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III. Claim 10, a pharmaceutical composition for vascular treatment. If this Group is elected, then the below Summarized Species Election is also required.

- IV. Claim 12, drawn to a therapeutic/prevention method for vascular disorders comprising administering the nuclear transfer promoter for Cdc42 protein in an effective amount for therapy/prevention to a patient in need of therapy/prevention of vascular disorders. If this Group is elected, then the below Summarized Species Election is also required.
- V. Claim 13-15, drawn to a method of screening a blood vessel remedy.
- VI. Claims 4, 5, 6, and 11, drawn to the "use" of a nuclear transfer promoter for Cdc42 protein.

The inventions represented above as Groups I-VI relate to a general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they share the same or corresponding technical features. Specifically, the technical feature of Groups I-VI is the nuclear transfer promoter for Cdc42 protein. The inventions lack unity, however, as the common technical feature is known in the art as evidenced by the teaching of Kohro et al. (WO 2005/079847 A1; abstract only) of a nuclear transfer promoter for cdc42 protein comprising an isoprenoid synthesis inhibitor and/or a geranylgeranyl transferase inhibitor (see abstract), and also Morikawa et al. (Morikawa et al. The effects of statins on mRNA levels of genes related to inflammation, coagulation, and vascular constriction in HUVEC, human umbilical vein endothelial cells. J. Atheroscler. Thromb. 2002; 9(4): 178-

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183; already made of record by applicant). Thus, the requirement is proper as the inventions represented above as Groups I-II lack unity of invention under PCT Rule 13.1.

Species Election regarding Groups I-VI

This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1. For example, the generic inventions encompass a multiplicity of chemically distinct isoprenoid synthesis inhibitors, geranylgeranyl transferase inhibitors, and various different combinations of a isoprenoid synthesis inhibitor and a geranylgeranyl transferase inhibitor, which would reasonably exhibit different/variable pharmacologic, pharmaceutical, therapeutic, side effects profiles. The therapeutic effects to be achieved with these different compounds or combination species would reasonably differ substantially depending on the specific compound, as well as the doses and duration of treatment, and the contemplated targeted vascular disorder intended. Each vascular disorder represents a different distinct clinical entity. Therapeutic modalities used to treat a certain vascular disorder may not be effective to treat other types of vascular disease. Thus, these species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single species for purposes of examination from the below list:

- 1a) a single chemically defined isoprenoid synthesis inhibitor e.g. pitavastatin; or
- 1b) a single chemically defined geranylgeranyl transferase inhibitor; or

1c) a single combination of a compound from above item "1a" and a compound from above item "1b."

Applicant is advised that a reply to this requirement <u>must include an identification of the</u>

species that is elected consonant with this requirement, and a listing of all claims readable

thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to the additional species which depend from or otherwise require all the limitations of an allowable generic claim as provided by 37 CFR 1.141. If claims are added after election, applicant <u>must</u> indicate which are readable upon the elected species (MPEP 809.02(a). Claims 1, 4, 5, 7, 10, 11, 12, and 13 are considered generic to the above species.

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and the product claims are subsequently found allowable, withdrawn process claims that depend from or otherwise require all the limitations of the allowable product claim will be considered for rejoinder. <u>All</u> claims directed to a nonelected process invention must require all the limitations of an allowable product claim for that process invention to be rejoined.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103 and 112. Until all claims to the elected product are found allowable, an otherwise proper

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restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowable product claim will not be rejoined. See MPEP § 821.04(b). Additionally, in order to retain the right to rejoinder in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution to require the limitations of the product claims. **Failure to do so**may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Inventorship Notice

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Charlesworth Rae whose telephone number is 571-272-6029. The examiner can normally be reached between 9 a.m. to 5:30 p.m. Monday to Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Woodward, can be reached at 571-272-8373. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR.

Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http:pair-direct.uspto.gov. Should you have any questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 800-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

10 May 2008 /C. R./ Examiner, Art Unit 1611

/Brian-Yong S Kwon/ Primary Examiner, Art Unit 1614